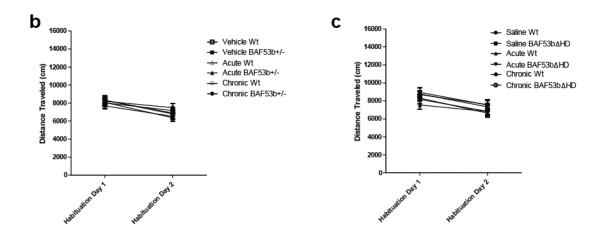
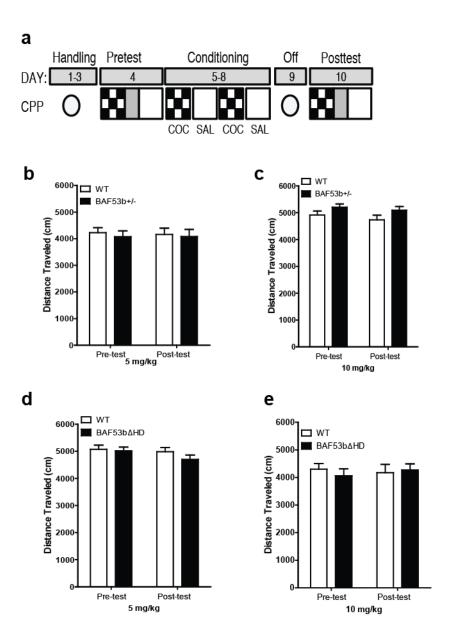
a	Handling	<u>Habituation</u>	Conditioning				
DAY:	1-3	4-5	6-10				
	0						
Contro	l		SAL	SAL	SAL	SAL	SAL
Acute			SAL	SAL	SAL	SAL	COC
Chroni	С		COC	COC	COC	COC	COC

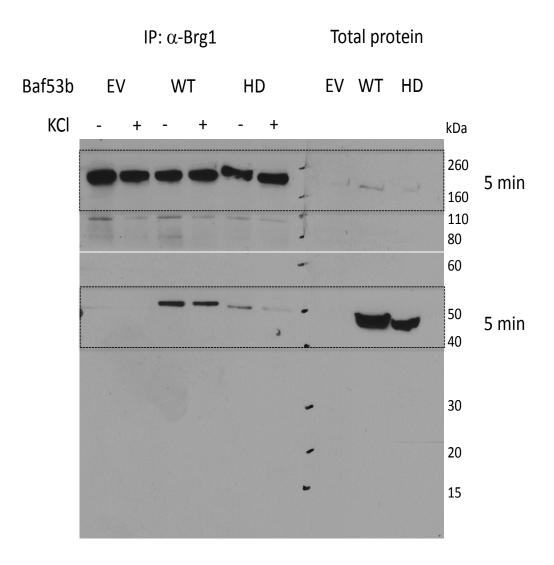


Supplementary Figure 1. BAF53b^{+/-} heterozygous knockout and BAF53bΔHD mice displayed normal baseline locomotor activity during habituation. (a) Schematic representation of cocaine sensitization procedure. (b) BAF53b^{+/-} heterozygous knockout and wildtype mice, regardless of future treatment, displayed similar locomotor activity throughout habituation (significant main effect of habituation day, $F_{1,24}$ =69.79, p<0.0001; no main effect of genotype, $F_{2,24}$ =0.41, p=0.67; no interaction, $F_{2,24}$ =0.91, p=0.41). (c) BAF53bΔHD and wild-type mice displayed similar locomotor activity throughout habituation (significant main effect of habituation day, $F_{1,20}$ =37.08, p<0.0001; no main effect of genotype, $F_{2,20}$ =2.02, p=0.16; no interaction, $F_{2,20}$ =1.38, p=0.27).



Supplementary Figure 2. Mutant mice have normal locomotion on test days during cocaine-CPP. **(a)** Schematic representation of cocaine-CPP procedure. **(b)** Cocaine-CPP expression indicated by mean CPP score (CS+ minus CS-) \pm S.E.M. At 5mg/kg cocaine dose, BAF53b^{+/-} heterozygous knockout mice (n=10) exhibited similar locomotion compared to wild-type littermates (n=8). A two-way repeated measures ANOVA revealed no main effect of genotype (F_{1,16}=0.14, p=0.72). No effect of conditioning was observed (F_{1,16}=0.09, p=0.77) and there was no interaction (F_{1,16}=0.16, p=0.69). **(c)** At 10mg/kg cocaine dose, BAF53b^{+/-} heterozygous knockout mice (n=9) exhibit similar CPP score to wild-type littermates (n=8). Using a two-way repeated measures ANOVA, we found no main effect of conditioning (F_{1,17}=2.22, p=0.15) nor genotype (F_{1,17}=3.07, p=0.09) and no interaction (F_{1,17}=0.09, p=0.77). **(d)** At 5mg/kg cocaine dose, BAF53bΔHD mice (n=9) exhibited significantly attenuated CPP score compared to wild-

type littermates (n=10). Using a two-way repeated measures ANOVA, we found significant main effects on conditioning ($F_{1,17}$ =5.20, p=0.03) but not genotype ($F_{1,17}$ = 0.70, p=0.42), and no interaction ($F_{1,17}$ =1.68, p=0.21). (e) At a 10mg/kg cocaine dose, BAF53b Δ HD mice (n=10) exhibited significantly attenuated CPP score compared to wild-type littermates (n=7).). A two-way repeated measures ANOVA revealed significant main effects of conditioning ($F_{1,15}$ =0.08, p=0.78), genotype ($F_{1,15}$ =0.05, p=0.83) and an interaction ($F_{1,15}$ =1.30, p=0.27).



Supplementary Figure 3. Immunoprecipitation and western blot showing that Brg1 (top dotted box) co-immunoprecipitates with BAF53b (Bottom dotted box) as well as BAF53b Δ HD. Image attained through 5 minute exposure.